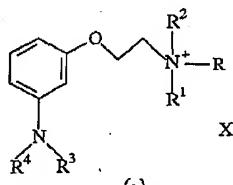


Appl. No. 10052,362
 Atty. Docket No. G-286M (CP-1218)
 Andt. Dated January 16th, 2004
 Reply to Office Action of October 9th, 2003
 Customer No. 27732

Amendments to the Specification

Please replace the paragraph beginning page 2 line 11, with the following paragraph:

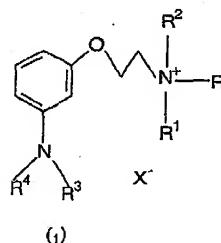
This invention provides novel couplers of the formula (1):



wherein X is selected from halogen and R⁵SO₄ where the halogen is preferably Cl, Br or I; R, R¹, and R² are each individually selected from C₁ to C₂₂ alkyl and C₁ to C₂₂ mono or dihydroxyalkyl, or two of R, R¹ and R² together with the nitrogen atom to which they are attached form a C₃ to C₆, preferably C₄ to C₆, saturated or unsaturated ring optionally containing in the ring one or more additional hetero atoms selected from O, S and N atoms; R³ and R⁴ are each individually selected from C₁ to C₆ alkyl, C₁ to C₆ hydroxyalkyl, C₁ to C₆ alkoxy, C₁ to C₆ aminoalkyl or R³ and R⁴ together form a C₄ to C₆ alkylene group; and R⁵ is selected from C₁ to C₂₂ alkyl and C₁ to C₂₂ mono or dihydroxyalkyl. Preferably X is Cl, Br, I and R⁵SO₄ where R⁵ is C₁ to C₄ alkyl, more preferably methyl; and preferably R, R¹, R², R³ and R⁴ are each individually C₁ to C₃ alkyl, and more preferably methyl.

Please replace the paragraph beginning page 3, line 10 with the following paragraph:

The coupler compounds of this invention are those of formula (1)

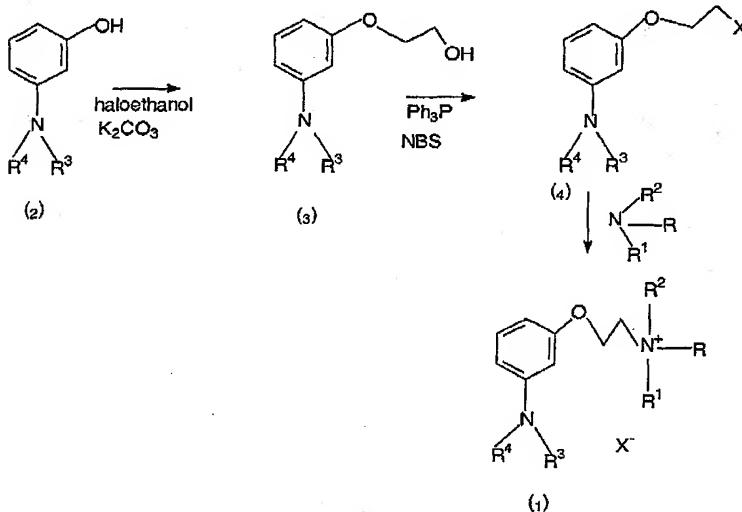


Appl. No. 10/052,362
 Atty. Docket No. G-286M (CP-1218)
 Audit. Dated January 16th, 2004
 Reply to Office Action of October 9th, 2003
 Customer No. 27732

wherein X is selected from halogen and R⁵SO₄ where the halogen is preferably Cl, Br or I; R, R¹, and R² are each individually selected from C₁ to C₂₂ alkyl and C₁ to C₂₂ mono or dihydroxyalkyl, or two of R, R¹ and R² together with the nitrogen atom to which they are attached form a C₃ to C₆, preferably C₄ to C₆, saturated or unsaturated ring optionally containing in the ring one or more additional hetero atoms selected from O, S and N atoms; R³ and R⁴ are each individually selected from C₁ to C₆ alkyl, C₁ to C₆ hydroxyalkyl, C₁ to C₆ alkoxy, C₁ to C₆ aminoalkyl or R³ and R⁴ together form a C₄ C₂ to C₆ alkylene group; and R⁵ is selected from C₁ to C₂₂ alkyl and C₁ to C₂₂ mono or dihydroxyalkyl. Preferably X is Cl, Br, I and R⁵SO₄ where R⁵ is C₁ to C₄ alkyl, more preferably methyl; and preferably R, R¹, R², R³ and R⁴ are each individually C₁ to C₃ alkyl, and more preferably methyl.

Please replace the paragraph beginning page 4, line 11 with the following paragraph:

The novel coupler compounds of formula (1) of this invention are readily prepared according to the following reaction sequence where X, R, R¹, R², R³, R⁴ and R⁵ are as defined hereinbefore:



Appl. No. 10/052,362
Atty. Docket No. G-286M (CP-1218)
Andt. Dated January 16th, 2004
Reply to Office Action of October 9th, 2003
Customer No. 27752

In this synthesis an aminophenol (2) is reacted with a 2-haloethanol, such as 2-bromoethanol, in the presence of potassium carbonate in dimethylformamide to produce the alcohol compound (3). Transformation of this alcohol compound (3) into a compound (4) is carried out by treatment of the alcohol compound with triphenylphosphine and a halo-succinimide, such as bromosuccinimide (NBS). Treatment of compound (4) with a quaternization reagent (NRR¹R²) produces a compounds of formula (1) of this invention.